WEST Search History

Hide Items Restore Clear Cancel

DATE: Friday, January 06, 2006

Hide?	Hide? Set Name Query Hi						
	DB=PGPB, USPT, EPAB; PLUR=YES; OP=ADJ						
	L12	L11 AND L9	84				
	L11	L10 AND L3	323				
	L10	L2 AND L1	1146				
	L9	L8 or L7	29257				
	L8	(435/7.1 435/7.23)![CCLS]	13124				
	L7	(530/350)![CCLS]	18276				
	L6	(530)![CCLS]	0				
	L5	(530)![CCLS]	0				
	L4	2A2A9	0				
	L3	prostate	38232				
	L2	tumor\$ or tumuor\$ or cancer\$ or neoplas\$	180591				
	L1	(jakobovits or afar or challita\$ or levin or mitchell or hubert).in.	34377				

END OF SEARCH HISTORY

GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

December 4, 2005, 09:57:24; Search time 187 Seconds Run on:

(without alignments)

1184.208 Million cell updates/sec

US-09-771-312-2 Title:

Perfect score: 2694

Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

2443163 segs, 439378781 residues Searched:

Total number of hits satisfying chosen parameters: 2443163

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_21:*

1: geneseqp1980s:*

2: geneseqp1990s:*

3: geneseqp2000s:*

4: geneseqp2001s:*

5: geneseqp2002s:*

6: geneseqp2003as:*

7: geneseqp2003bs:*

8: geneseqp2004s:*

9: geneseqp2005s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	<pre>% Query Match</pre>	Length	DB	ID	Description	
1	2694	100.0	504	4	AAU06524	Aau06524 Prostate	2
2	2694	100.0	528	4	AAB92632	Aab92632 Human pr	ro
3	2694	100.0	528	5	ABB97288	Abb97288 Novel hi	um
4	1808	67.1	376	8	ADR99239	Adr99239 Hypothet	ti
5	825.5	30.6	313	4	ABG08002	Abg08002 Novel hu	am
6	608.5	22.6	423	4	ABG23408	Abg23408 Novel hu	am
7	591.5	22.0	446	5	ABB75706	Abb75706 Human ph	no
8	588.5	21.8	482	5	ABP43772	Abp43772 14 clone	2

```
9
    533.5
            19.8
                    453 5 ABB97561
                                                    Abb97561 Novel hum
                    351 4 AAB94662
10
    530.5
            19.7
                                                    Aab94662 Human pro
    530.5
            19.7
11
                    351 5
                           ABB97470
                                                    Abb97470 Novel hum
            12.5
12
      338
                    275
                           AAB92468
                                                    Aab92468 Human pro
13
    320.5 11.9
                    223 4
                           AAM15386
                                                   Aam15386 Peptide #
14
    320.5 11.9
                    223 4 ABB34392
                                                   Abb34392 Peptide #
15
   320.5 11.9
                    223 4 AAM27874
                                                   Aam27874 Peptide #
16
   320.5 11.9
                    223 4 ABB29229
                                                    Abb29229 Peptide #
17
    320.5
           11.9
                    223 4 AAM67577
                                                   Aam67577 Human bon
18
    320.5
            11.9
                    223 4
                           AAM55182
                                                  Aam55182 Human bra
19
    320.5
           11.9
                   223 4 ABG49223
                                                  Abg49223 Human liv
20
    320.5
           11.9
                    223 4 AAM03148
                                                  Aam03148 Peptide #
21
    320.5 11.9
                   223 5 ABG37168
                                                   Abg37168 Human pep
22
    320.5
           11.9
                   223 8 ABO59933
                                                    Abo59933 Human gen
23
    236.5
            8.8
                   123 4
                           AAM95677
                                                    Aam95677 Human rep
24
      196
             7.3
                   1038 7
                           ADC03412
                                                    Adc03412 Rice flow
25
      196
             7.3
                   1038
                        7
                           ABM88777
                                                    Abm88777 Rice abio
26
    181.5
             6.7
                   564 8
                           ADY23792
                                                    Ady23792 Plant ful
27
    167.5
             6.2
                   767 6 ABR53431
                                                   Abr53431 Protein s
    167.5
28
             6.2
                   767 7 ADK64670
                                                   Adk64670 Disease t
29
    164.5
                    554 3 AAG36165
             6.1
                                                   Aag36165 Arabidops
30
    164.5
                    652 3 AAG36164
             6.1
                                                    Aaq36164 Arabidops
                   781 3 AAG36163
31
    164.5
             6.1
                                                    Aaq36163 Arabidops
                    815 5 AAG78388
32
    162.5
             6.0
                                                   Aag78388 Human H37
33
    162.5
             6.0
                   815 7
                           AAE38620
                                                  Aae38620 Human H37
    162.5
34
             6.0
                   815 7 AAE38621
                                                  Aae38621 Human H37
35
    162.5
             6.0
                    815 8 ADP23184
                                                  Adp23184 PRO polyp
36
    162.5
             6.0
                    815 9 ADX05546
                                                  Adx05546 Cyclin-de
37
      162
             6.0
                    381 2 AAY07056
                                                    Aay07056 Renal can
38
    157.5
             5.8
                    852
                        7
                           ADD45318
                                                    Add45318 Rat Prote
39
    157.5
             5.8
                    852
                        7
                           ADE56352
                                                   Ade56352 Rat Prote
    156.5
40
             5.8
                   573
                           ADM19760
                                                   Adm19760 Protein e
41
      156
             5.8
                    852 7
                           AEA79131
                                                  Aea79131 Human apo
42
      156
             5.8
                    852 9 ADX07612
                                                  Adx07612 Cyclin-de
43
                    929 4 AAM78604
      156
             5.8
                                                   Aam78604 Human pro
44
      156
                    930 8 ABM82400
             5.8
                                                   Abm82400 Tumour-as
                    930 9 ADX07610
45
      156
             5.8
                                                    Adx07610 Cyclin-de
```

ALIGNMENTS

```
RESULT 1
AAU06524
     AAU06524 standard; protein; 504 AA.
XX
AC
     AAU06524;
XX
DΤ
     24-OCT-2001 (first entry)
XX
     Prostate and testis-related gene 84P2A9 encoded protein.
DΕ
XX
KW
     84P2A9-related protein; prostate; testis; tissue; cancer; leukaemia;
KW
     tumour; kidney; brain; bone; skin; ovary; breast; pancreas; colon; lung;
KW
     cytostatic; gene therapy; antibody therapy; ribozyme; serum; blood;
KW
     single chain monoclonal antibody; urine.
XX
```

```
OS
    Homo sapiens.
XX
PN
    WO200155391-A2.
XX
PD
    02-AUG-2001.
XX
ΡF
    26-JAN-2001; 2001WO-US002651.
XX
    26-JAN-2000; 2000US-0178560P.
PR
XX
PΑ
    (UROG-) UROGENESYS INC.
XX
    Jakobovits A, Afar DEH, Challita-Eid PM, Levin E, Mitchell SC;
PΙ
PΙ
    Hubert RS;
XX
DR
    WPI; 2001-502631/55.
    N-PSDB; AAS11663.
DR
XX
PT
    New 84P2A9 gene and its encoded protein, useful for diagnosing and
    treating cancer, e.g. leukemia and cancer of the prostate, testis,
PT
PT
    kidney, brain or bone, or for eliciting an immune response.
XX
PS
    Claim 13; Fig 2; 149pp; English.
XX
    The polypeptide sequences represent the 84P2A9-related protein and
CC
CC
    peptide fragments of the protein. 84P2A9 exhibits prostate and testis
    specific expression in normal adult tissue, but it is also aberrantly
CC
CC
    expressed in many cancers including leukaemia and tumours of the
    prostate, testis, kidney, brain, bone, skin, ovary, breast, pancreas,
CC
CC
    colon and lung. The 84P2A9 polynucleotide, its related protein and
CC
    peptide fragments and specific PCR primers are therefore useful for
CC
    diagnosing and treating cancer. A vector comprising a polynucleotide
CC
    which encodes a single chain monoclonal antibody, that immunospecifically
    binds to an 84P2A9-related protein, and a ribozyme capable of cleaving a
CC
CC
    polynucleotide having the 84P2A9 coding sequence, are both useful in the
CC
    preparation of a composition for treating a patient with a cancer that
    expresses 84P2A9. The sequences can be used in diagnostic methods to
CC
CC
    monitor the level of 84P2A9 gene products in serum, blood, urine and
CC
    tissue and to thereby detect the presence of cancerous cells
XX
SO
    Sequence 504 AA;
                        100.0%; Score 2694; DB 4; Length 504;
 Query Match
 Best Local Similarity
                        100.0%; Pred. No. 3.9e-228;
 Matches 504; Conservative
                            0; Mismatches
                                               0; Indels
                                                            0; Gaps
                                                                       0;
           1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
QУ
             1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Db
Qу
          61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
             61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Db
Qу
         121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
             121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Db
```

```
181 TKNKVKKRKLKIIROGPKIODEGVVLESEETNOTNKDKMECEEOKVSDELMSESDSSSLS 240
Ov
           181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Db
        241 STDAGLFTNDEGROGDDEOSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Qу
           241 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Db
        301 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
QУ
           301 ILTGSFPLMSHPSRRGFOARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
Db
        361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Qу
           361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Db
        421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Οv
           421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Db
        481 GLGLGFPLPKSTSATTTPNAGKSA 504
Qу
           Db
        481 GLGLGFPLPKSTSATTTPNAGKSA 504
RESULT 2
AAB92632
ID
    AAB92632 standard; protein; 528 AA.
XX
AC
    AAB92632;
XX
DT
    26-JUN-2001 (first entry)
XX
    Human protein sequence SEQ ID NO:10938.
DΕ
XX
KW
    Human; primer; detection; diagnosis; antisense therapy; gene therapy.
XX
OS
    Homo sapiens.
XX
ΡN
    EP1074617-A2.
XX
    07-FEB-2001.
PD
XX
    28-JUL-2000; 2000EP-00116126.
PF
XX
    29-JUL-1999;
                99JP-00248036.
PR
                99JP-00300253.
PR
    27-AUG-1999;
    11-JAN-2000; 2000JP-00118776.
PR
    02-MAY-2000; 2000JP-00183767.
PR
    09-JUN-2000; 2000JP-00241899.
PR
XX
    (HELI-) HELIX RES INST.
PΑ
XX
    Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PΙ
PΙ
    Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
```

DR WPI; 2001-318749/34. XX PΤ Primer sets for synthesizing polynucleotides, particularly the 5602 full-PΤ length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length PΤ PTcDNAs. XX Claim 8; SEQ ID NO 10938; 2537pp + Sequence Listing; English. PS XX CC The present invention describes primer sets for synthesising 5602 full-CC length cDNAs defined in the specification. Where a primer set comprises: CC (a) an oligo-dT primer and an oligonucleotide complementary to the CC complementary strand of a polynucleotide which comprises one of the 5602 CC nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination CC CC of an oligonucleotide comprising a sequence complementary to the CC complementary strand of a polynucleotide which comprises a 5'-end CC sequence and an oligonucleotide comprising a sequence complementary to a CC polynucleotide which comprises a 3'-end sequence, where the CC oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the CC CC specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, CC particularly full-length cDNAs. The primers are also useful for the CC CC detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length CC CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 CC CC represent human amino acid sequences; and AAH13629 to AAH13632 represent CC oligonucleotides, all of which are used in the exemplification of the CC present invention XX SO Sequence 528 AA; 100.0%; Score 2694; DB 4; Query Match Length 528; Best Local Similarity 100.0%; Pred. No. 4.2e-228; Matches 504; Conservative 0; Mismatches 0: Indels 0; Gaps 0: Qy 1 MEELVHDLVSALEESSEOARGGFAETGDHSRSISCPLKROARKRRGRKRRSYNVHHPWET 60 Db 25 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 84 61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDOMLVAKRRPSSNLNNNVRGKRPLWH 120 Qу 85 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 144 Db 121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180 Qу 145 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 204 Db 181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240 Qу

> 241 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300

265 STDAGLFTNDEGROGDDEOSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 324

Db

Qγ

Db

```
301 ILTGSFPLMSHPSRRGFOARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
Qу
            Db
         325 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 384
         361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Qу
            Db
         385 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 444
         421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Qу
            445 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 504
Db
         481 GLGLGFPLPKSTSATTTPNAGKSA 504
Qу
            Db
         505 GLGLGFPLPKSTSATTTPNAGKSA 528
RESULT 3
ABB97288
    ABB97288 standard; protein; 528 AA.
XX
AC
    ABB97288;
XX
DΤ
    28-JUN-2002 (first entry)
XX
DΕ
    Novel human protein SEQ ID NO: 556.
XX
KW
    Human; antianaemic; vulnerary; antiinflammatory; immunomodulator;
    antiinfertility; cerebroprotective; cytostatic; rheumatic; gene therapy;
KW
KW
    neuroprotective; antiparkinsonian; protein therapy; EST;
KW
    expressed sequence tag.
XX
OS
    Homo sapiens.
XX
PN
    W0200222660-A2.
XX
PD
    21-MAR-2002.
XX
ΡF
    10-SEP-2001; 2001WO-US026015.
XX
PR
    11-SEP-2000; 2000US-00659671.
XX
PΑ
    (HYSE-) HYSEQ INC.
XX
PΙ
    Tang YT, Liu C,
                    Zhou P, Asundi V, Zhang J, Zhao QA, Ren F;
PΙ
    Xue AJ, Yang Y,
                    Wehrman T, Drmanac RT;
XX
DR
    WPI; 2002-292408/33.
    N-PSDB; ABN32474.
DR
XX
    An isolated polynucleotide for treating diseases associated with its
PT
PT
    encoded polypeptide such as cancer and multiple sclerosis.
XX
PS
    Example 2; SEQ ID NO 556; 509pp; English.
XX
    The present invention provides the protein and coding sequences of 444
CC
```

```
CC
    novel human proteins. These were isolated from expressed sequences tags
    (ESTs). They can be used to stimulate cell growth, to regulate
CC
CC
    haematopoiesis e.g. to treat aplastic anaemia, to help tissue regrowth
CC
    e.g. in burn treatment, to regulate the immune system e.g. to treat
CC
    multiple sclerosis, to regulate activin or inhibin e.g. to treat
CC
    infertility, to regulate haemostasis or thrombolysis e.g. to treat stroke
CC
    and cancer, to screen for drugs, to treat inflammatory conditions e.g.
CC
    rheumatoid arthritis, and to treat nervous system disorders e.g.
    Parkinson's disease. The present sequence is a protein of the invention
CC
XX
SQ
    Sequence 528 AA;
                     100.0%; Score 2694; DB 5;
 Query Match
                                            Length 528;
 Best Local Similarity
                     100.0%; Pred. No. 4.2e-228;
 Matches 504; Conservative
                         0; Mismatches
                                         0;
                                            Indels
                                                    0;
                                                        Gaps
                                                              0;
         1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Qу
           25 MEELVHDLVSALEESSEOARGGFAETGDHSRSISCPLKROARKRRGRKRRSYNVHHPWET 84
Db
         61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Qу
           85 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDOMLVAKRRPSSNLNNNVRGKRPLWH 144
Db
        121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPODISNKRTMTOPPEGCRDODMDSDRAYOYOEF 180
Qv
           145 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 204
Db
        181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Qу
           205 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 264
Db
        241 STDAGLFTNDEGROGDDEOSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Qу
           265 STDAGLFTNDEGROGDDEOSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 324
Db
        301 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 360
Qу
           325 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSMVPIPGPVGNKRMVHFSPD 384
Db
        361 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Qу
           Db
        385 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 444
        421 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Qу
           Db
        445 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 504
        481 GLGLGFPLPKSTSATTTPNAGKSA 504
Qу
           Db
        505 GLGLGFPLPKSTSATTTPNAGKSA 528
```

RESULT 4 ADR99239

ID ADR99239 standard; protein; 376 AA.

XX

```
АC
    ADR99239;
XX
DT
     02-DEC-2004 (first entry)
XX
     Hypothetical protein FLJ10252, SEQ ID 245.
DE
XX
KW
     Cytostatic; breast cancer; cancer; human; FLJ10252.
XX
OS
    Homo sapiens.
XX
PN
    WO2004078035-A2.
XX
     16-SEP-2004.
PD
XX
PF
     27-FEB-2004; 2004WO-US007268.
XX
     28-FEB-2003; 2003US-0450655P.
PR
XX
     (FARB ) BAYER PHARM CORP.
PA
XX
PΙ
     Eveleigh D, Bigwood D;
XX
     WPI; 2004-653556/63.
DR
DR
     N-PSDB; ADR99112.
XX
     Diagnosing breast cancer comprises comparing the level of expression of
PΤ
РΤ
     genes or gene products in a first biological sample taken from a patient
PT
     with that in a normal patient sample.
XX
PS
     Claim 3; SEQ ID NO 245; 53pp; English.
XX
     The present invention relates to a method (M1) for diagnosing breast
CC
     cancer in a patient. The method comprises comparing the level of
CC
     expression of one or more genes or gene products in a biological sample
CC
CC
     from the patient with that in a normal patient sample, where a difference
CC
     in the gene expression in the first sample compared to that in the second
CC
     sample is a diagnostic of the disease. Also claimed are: method (M2) for
     distinguishing between normal and disease tissues; method (M3) for
CC
CC
     monitoring the response of a breast cancer patient to treatment with an
CC
     anti-cancer agent; method (M4) for identifying a compound for treating
CC
     breast cancer; and an array for distinguishing between normal and disease
CC
     tissues comprising two or more probes corresponding to genes selected
CC
     from ADR98995-ADR99121 or comprising two or more polypeptides selected
CC
     from ADR99122-ADR99248. In M1 and M2 the genes are selected from ADR98995
     -ADR99121 and the gene products are polypeptides selected from ADR99122-
CC
     ADR99248. M1 is useful for diagnosing breast cancer. M2 and the array are
CC
CC
     useful for distinguishing between normal and disease tissue. M3 is useful
CC
     for monitoring the response of a breast cancer patient to treatment with
CC
     an anti-cancer agent. M4 is useful for identifying a compound for
     treating breast cancer. Note: The sequence data for this patent did not
CC
     form part of the printed specification, but was obtained in electronic
CC
     format directly from WIPO at ftp.wipo.int/pub/published pct sequences.
CC
XX
SQ
     Sequence 376 AA;
```

67.1%; Score 1808; DB 8; Length 376;

Best Local Similarity 99.7%; Pred. No. 3.1e-150;

Query Match

```
1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Qу
           25 MEELVHDLVSALEESSEOARGGFAETGDHSRSISCPLKROARKRRGRKRRSYNVHHPWET 84
Db
         61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Qу
           85 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 144
Db
        121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Qy
           145 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 204
Db
        181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
Qу
           205 TKNKVKKRKLKIIRQGPKIQNEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 264
Db
        241 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 300
Qу
           265 STDAGLFTNDEGRQGDDEQSDWFYEKESGGACGITGVVPWWEKEDPTELDKNVPDPVFES 324
Db
        301 ILTGSFPLMSHPSRRGFQARLSRLHGMSSKNIKKSGGTPTSM 342
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KW
    Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW
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PA
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    Drmanac RT, Liu C, Tang YT;
XX
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DR
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0; Indels

0;

Gaps

0;

Matches 341; Conservative 1; Mismatches

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PT
    New isolated polynucleotide and encoded polypeptides, useful in
    diagnostics, forensics, gene mapping, identification of mutations
PT
    responsible for genetic disorders or other traits and to assess
PΤ
PT
    biodiversity.
XX
PS
    Claim 20; SEQ ID NO 38361; 103pp; English.
XX
    The invention relates to isolated polynucleotide (I) and polypeptide (II)
CC
CC
    sequences. (I) is useful as hybridisation probes, polymerase chain
CC
    reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
CC
    and in recombinant production of (II). The polynucleotides are also used
CC
    in diagnostics as expressed sequence tags for identifying expressed
CC
    genes. (I) is useful in gene therapy techniques to restore normal
    activity of (II) or to treat disease states involving (II). (II) is
CC
    useful for generating antibodies against it, detecting or quantitating a
CC
    polypeptide in tissue, as molecular weight markers and as a food
CC
    supplement. (II) and its binding partners are useful in medical imaging
CC
    of sites expressing (II). (I) and (II) are useful for treating disorders
CC
    involving aberrant protein expression or biological activity. The
CC
    polypeptide and polynucleotide sequences have applications in
CC
    diagnostics, forensics, gene mapping, identification of mutations
CC
    responsible for genetic disorders or other traits to assess biodiversity
CC
    and to produce other types of data and products dependent on DNA and
CC
    amino acid sequences. ABG00010-ABG30377 represent novel human diagnostic
CC
CC
    amino acid sequences of the invention. Note: The sequence data for this
    patent did not appear in the printed specification, but was obtained in
CC
    electronic format directly from WIPO at
CC
CC
    ftp.wipo.int/pub/published pct sequences
XX
SQ
    Sequence 313 AA;
                      30.6%; Score 825.5; DB 4; Length 313;
  Query Match
  Best Local Similarity 55.7%; Pred. No. 9.2e-64;
  Matches 181; Conservative 4; Mismatches 25; Indels 115; Gaps
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Qу
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Qу
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296 PVFESILTGSFPLMSHPSRRGFQAR 320

Qу

GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame plus p2n model Run on: December 11, 2005, 17:36:59; Search time 7575 Seconds (without alignments) 3782.059 Million cell updates/sec US-09-771-312-2 Title: Perfect score: 2694 Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504 Scoring table: BLOSUM62 Xgapop 10.0 , Xgapext Ygapop 10.0 , Ygapext 0.5 Fgapop 6.0 , Fgapext 7.0 Delop 6.0 , Delext Searched: 5883141 segs, 28421725653 residues Total number of hits satisfying chosen parameters: 11766282 Minimum DB seq length: 0 Maximum DB seq length: 2000000000 Post-processing: Minimum Match 0% Maximum Match 100% Listing first 45 summaries Command line parameters: -MODEL=frame+ p2n.model -DEV=xlh -Q=/cgn2 1/USPTO spool/US09771312/runat 01122005 145311 15042/app query.fasta 1.647 -DB=GenEmbl -QFMT=fastap -SUFFIX=rge -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pto -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000 -USER=US09771312_@CGN_1_1_4939_@runat_01122005_145311_15042 -NCPU=6 -ICPU=3 -NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG -DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7 Database : GenEmbl:* 1: gb ba:* 2: gb_in:* 3: gb env:* 4: gb om:* 5: gb ov:* 6: gb_pat:*

7: gb_ph:*
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9: gb_ro:*
10: gb sts:*

11: gb_sy:*
12: gb_un:*
13: gb_vi:*
14: gb_htg:*
15: gb pl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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AK001114 Homo sapi
AX405697 Sequence
AX206855 Sequence
BC054810 Mus muscu
AJ851518 Gallus ga
CQ720787 Sequence
BD183390 Novel gen
BC042193 Homo sapi
BC063474 Homo sapi
BC097745 Xenopus l
BC079232 Rattus no
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AK129299 Mus muscu
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BX004824 Zebrafish 2694 100.0 2338 6 AX876032 AX876032 Sequence 2694 100.0 2338 8 AK001114 3 2694 100.0 2344 6 AX405697 4 2694 100.0 2345 6 AX206855

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DEFINITION Primer for synthesizing full-length cDNA and use thereof.
ACCESSION
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VERSION
            BD155908.1 GI:27861666
            JP 2002191363-A/10751.
KEYWORDS
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            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE
            1 (bases 1 to 2338)
  AUTHORS
            Ota, T., Isogai, T., Nishikawa, T., Hayashi, K., Saito, K.,
Yamamoto, J.,
            Ishii, S., Sugiyama, T., Wakamatsu, A., Nagai, K. and Otsuki, T.
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            Patent: JP 2002191363-A 10751 09-JUL-2002;
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            HELIX RESEARCH INSTITUTE
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            PΙ
                 SAITO,
            PΙ
                 JUNICHI YAMAMOTO, SHIZUKO ISHII, TOMOYASU SUGIYAMA, AI
WAKAMATSU,
            PΙ
                 KEIICHI NAGAI, TETSUJI OTSUKI
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Alignment Scores:

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Query Match: 100.00% Indels: 0
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QУ	381	LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
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2003
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VERSION
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            Hominidae; Homo.
REFERENCE
  AUTHORS
            Ota, T., Isogai, T., Nishikawa, T., Hayashi, K., Saito, K.,
Yamamoto, J.,
            Ishii, S., Sugiyama, T., Wakamatsu, A., Nagai, K. and Otsuki, T.
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            Primers for synthesising full-length cDNA and their use
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ORIGIN

Alignment Score: Percent Simin Best Local Souery Match: DB:	larity:	5.58e-138 2694.00 100.00% 100.00% 100.00%	Length: Matches: Conservative: Mismatches: Indels: Gaps:	2338 504 0 0 0	
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Qy 4				ValHisHisProTrpGluThr	60
Db 29					350
О У 6				GluProSerLysAspTyrArg	80
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Qy 8				AspAspGlnMetLeuValAla	100
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Qу
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2004
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ACCESSION
           AK001114
VERSION
           AK001114.1 GI:7022173
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SOURCE
           Homo sapiens (human)
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            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
 AUTHORS
           Ota, T., Suzuki, Y., Nishikawa, T., Otsuki, T., Sugiyama, T., Irie, R.,
           Wakamatsu, A., Hayashi, K., Sato, H., Nagai, K., Kimura, K., Makita, H.,
            Sekine, M., Obayashi, M., Nishi, T., Shibahara, T., Tanaka, T.,
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            Kumagai, A., Itakura, S., Fukuzumi, Y., Fujimori, Y., Komiyama, M.,
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Kobatake, N.,
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            Nagase, T., Nomura, N., Kikuchi, H., Masuho, Y., Yamashita, R.,
            Nakai, K., Yada, T., Nakamura, Y., Ohara, O., Isogai, T. and Sugano, S.
  TITLE
            Complete sequencing and characterization of 21,243 full-length
            human cDNAs
  JOURNAL
            Nat. Genet. 36 (1), 40-45 (2004)
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   PUBMED
REFERENCE
            Isogai, T., Ota, T., Hayashi, K., Sugiyama, T., Otsuki, T., Suzuki, Y.,
 AUTHORS
            Nishikawa, T., Nagai, K., Sugano, S., Shiratori, A., Sudo, H.,
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  TITLE
            NEDO human cDNA sequencing project
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               (bases 1 to 2338)
REFERENCE
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 AUTHORS
            Isogai, T. and Otsuki, T.
  TITLE
            Direct Submission
  JOURNAL
            Submitted (16-FEB-2000) Takao Isogai, Helix Research Institute,
            Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
            (E-mail:genomics@hri.co.jp, Tel:81-438-52-3975, Fax:81-438-52-
3986)
COMMENT
            NEDO human cDNA sequencing project supported by Ministry of
            International Trade and Industry of Japan; cDNA full insert
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ORIGIN

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US-09-771-312-2 (1-504) x AK001114 (1-2338)

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2002
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VERSION
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REFERENCE
          Tang, Y.T., Liu, C., Zhou, P., Asundi, V., Zhang, J., Zhao, Q.A.,
  AUTHORS
Ren, F.,
          Xue, A.J., Yang, Y., Wehrman, T. and Drmanac, R.T.
          Novel nucleic acids and polypeptides
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ORIGIN

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	REFERENCE AUTHORS	: 1 : Ja	akobovits, A., Afar, D.E., Challita-Eid, P.M., Levin, E., itchell, S.C. and Hubert, R.S.					

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84p2a9: a prostate and testis specific protein highly expressed in
 TITLE
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 JOURNAL
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          Urogenesys, Inc. (US)
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Qу	201	$Asp {\tt GluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu}$	220
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DEFINITION Mus musculus G patch domain containing 2, mRNA (cDNA clone MGC:65681 IMAGE:6839419), complete cds.						
VERSION		BC054810 BC054810.1 GI:32452009 MGC.				

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ORGANISM Mus musculus
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REFERENCE
                (bases 1 to 4537)
  AUTHORS
            Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
            Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M.,
Schuler, G.D.,
            Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
            Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
            Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
            Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
            Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,
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  TITLE
            Generation and initial analysis of more than 15,000 full-length
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            Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
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            Strausberg, R.
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  JOURNAL
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            Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
            NIH-MGC Project URL: http://mgc.nci.nih.gov
  REMARK
COMMENT
            Contact: MGC help desk
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Dr. James Lin, University of Iowa
            cDNA Library Preparation: M. Bento Soares, University of Iowa
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: University of Iowa, Dr. M. Bento Soares and Dr.
            Thomas L. Casavant.
            Web site: http://genome.uiowa.edu
            Contact: bento-soares@uiowa.edu; tom-casavant@uiowa.edu
            Bonaldo, M.F., Akabogu, I., Bair, T., Bair, J., Crouch, K., Davis, A.,
            Fishler, K., Keppel, C., Kucaba, T., Lebeck, M., Melo, A., Schaefer, K.,
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Location/Qualifiers

SOURCE

FEATURES

Mus musculus (house mouse)

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ORIGIN

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Score:
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US-09-771-312-2 (1-504) x BC054810 (1-4537)
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Db 1174	1115	CTCAGTCGCCTTCATGGAACGCCTTCAAAGAATATTAAAAAGTCTTCAGGGGCTCCACCT	
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us-09-771-312-2.rng

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11: 12:

13: 14:

SUMMARIES

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PΙ
XX
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DR
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        Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length
PT
PT
PT
PΤ
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XX
         Claim 8; SEQ ID NO 10937; 2537pp + Sequence Listing; English.
PS
XX
        The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602
CC
CC
CC
CC
        nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC
CC
        of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end
CC
\mathsf{CC}
        sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the
CC
CC
CC
CC
        specification. The primer sets can be used in antisense therapy and in
CC
        gene therapy. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC
CC
CC
CC
CC
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CC
CC
         represent human amino acid sequences; and AAH13629 to AAH13632 represent
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CC
         present invention
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US-09-771-312-2 (1-504) x AAH13916 (1-2338)

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Db		GÁGTCTGÁTTTTGCTGTGGACAATGTTGGGAATAGAACTCTGCGCAGGAGGAGAAAGGTA 590	
Qy		LysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMetThrGlnPro	
Db		ÁÁÁCGCÁTGGCÁGTÁGÁTCTCCCACÁGGACATCTCTAACAAACGGACAATGACCCAGCCA 650	
Qy		ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe 180	
Db		CCTGAGGGTTGTAGAGATCAGGACATGGACAGTGATAGAGCCTACCAGTATCAAGAATTT 710	
Qy		ThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyProLysIleGln 200	
Db		ACCAAGAACAAAGTCAAAAAAAAAAGAAAGTTGAAAATAATCAGACAAGGACCAAAAATCCAA 770	
Qy		AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu 220	
Db		GATGAAGGAGTAGTTTTAGAAAGTGAGGAAACGAACCAGACCAATAAGGACAAAATGGAA 830	
Qy		CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer 240	
Db		TGTGAAGAGCAAAAAGTCTCAGATGAGCTCATGAGTGAAAGTGATTCCAGCAGTCTCAGC 890	
Qy		SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer 260	
Db		AĞCACTĞATGCTĞĞATTĞTTTACCAATGATGAGGGAAGACAAGGTGATGAACAGAGT 950 ASpTrpPhetyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp 280	
Qy Db			
Qy		TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer 300	
Dp		TIPETULYSGIUASPITOTIII GIULEUASPLYSASIIVUITTOASPITOVUITTILGIUSET 300 IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	
55	1011	Danie 4	

```
301 IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg 320
Qy
        1071 ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA 1130
Db
         321 LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr 340
Qy
Db
        1131 CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAGGGACTCCAACT 1190
         341 SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp 360
Qy
        1191 TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTTCCCCGGAT 1250
Db
         361 SerHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln 380
Qy
        1251 TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG 1310
Db
         381 LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
Qy
        1311 CTTCTGAGAGATAATCGAGCTGAAAGAGGACACAAGAAAAATTGTTCTGTGAGAACAGCC 1370
Db
         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
Db
        1371 ÁGCÁGGCÁÁÁCÁÁGCÁTGCÁTTTÁGGÁTCCTTÁTGCÁCGGÁGÁTÁTCÁÁÁCGGÁGÁÁÁGÁ 1430
         421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
             1431 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1490
Db
         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
             Db
        1491 ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA 1550
         461 GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys 480
Qy
        1551 GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG 1610
Db
         481 GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
             Db
        1611 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1670
         501 GlyLysSerAla 504
Qy
        1671 GGAAAATCCGCC 1682
Db
RESULT 2
ADR99112
    ADR99112 standard; DNA; 2338 BP.
ID
XX
AC
    ADR99112;
XX
    02-DEC-2004 (first entry)
DT
XX
    Hypothetical protein FLJ10252, coding sequence, SEQ ID 118.
DE
XX
    Cytostatic; breast cancer; cancer; human; gene; ds; FLJ10252.
KW
XX
os
    Homo sapiens.
XX
    WO2004078035-A2.
PΝ
XX
    16-SEP-2004.
PD
XX
```

```
us-09-771-312-2.rng
PF
       27-FEB-2004; 2004WO-US007268.
XX
PR
       28-FEB-2003: 2003US-0450655P.
XX
       (FARB ) BAYER PHARM CORP.
PA
XX
PΙ
       Eveleigh D, Bigwood D;
XX
       WPI; 2004-653556/63.
DR
       P-PSDB; ADR99239.
REFSEQ; NM_018040.1.
DR
DR
XX
       Diagnosing breast cancer comprises comparing the level of expression of genes or gene products in a first biological sample taken from a patient
PT
PT
PT
       with that in a normal patient sample.
XX
PS
       Claim 2; SEQ ID NO 118; 53pp; English.
XX
       The present invention relates to a method (M1) for diagnosing breast
CC
       cancer in a patient. The method comprises comparing the level of expression of one or more genes or gene products in a biological sample
CC
CC
       from the patient with that in a normal patient sample, where a difference
CC
CC
       in the gene expression in the first sample compared to that in the second
       sample is a diagnostic of the disease. Also claimed are: method (M2) for
CC
       distinguishing between normal and disease tissues; method (M3) for
CC
      monitoring the response of a breast cancer patient to treatment with an anti-cancer agent; method (M4) for identifying a compound for treating breast cancer; and an array for distinguishing between normal and disease tissues comprising two or more probes corresponding to genes selected from ADR98995-ADR99121 or comprising two or more polypeptides selected from ADR99122-ADR99248. In M1 and M2 the genes are selected from ADR98995
CC
CC
CC
CC
CC
CC
       -ADR99121 and the gene products are polypeptides selected from ADR99122-ADR99248. M1 is useful for diagnosing breast cancer. M2 and the array are useful for distinguishing between normal and disease tissue. M3 is useful for monitoring the response of a breast cancer patient to treatment with an anti-cancer agent. M4 is useful for identifying a compound for
CC
CC
CC
CC
CC
       treating breast cancer. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic
CC
CC
CC
       format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
       Sequence 2338 BP; 739 A; 478 C; 541 G; 580 T; 0 U; 0 Other;
Alignment Scores:
Pred. No.:
                                  2.07e-199
                                                                             2338
                                                       Length:
Score:
                                  2694.00
                                                       Matches:
                                                                             504
Percent Similarity:
                                  100.00%
                                                        Conservative:
                                                                             0
Best Local Similarity:
                                  100.00%
                                                                             0
                                                       Mismatches:
Query Match:
                                  100.00%
                                                       Indels:
                                                                             0
                                                                             0
US-09-771-312-2 (1-504) x ADR99112 (1-2338)
Qy
                 1 MetGluGluLeuValHisAspLeuValSerAlaLeuGluGluSerSerGluGlnAlaArg 20
                    Db
              171 ÁTGGÁGGÁGCTGGTTCÁTGÁCCTTGTCTCAGCATTGGAAGAGAGCTCAGAGCAAGCTCGA 230
               21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40
Qy
                     Db
              231 GGTGGATTTGCTGAAACAGGAGACCATTCTCGAAGTATATCTTGCCCTCTGAAACGCCAG 290
               41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60
Qy
```

291 GCÁAGGÁAÁAGGAGAGGAGAAAACGGÁGGTCGTÁTÁÁTGTGCÁTCACCCGTGGGAGACT 350

Db

Qy	61	GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg	80
Db	351	GGTCACTGCTTAAGTGAAGGCTCTGATTCTAGTTTAGAAGAACCAAGCAAG	410
Qy	81	GluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMetLeuValAla	100
Db	411	GAGAATCACAATAATAATAAAAAAAGATCACAGTGACTCTGATGACCAAATGTTAGTAGCA	470
Qy	101	LysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgProLeuTrpHis	120
Db	471	AAGCGCAGGCCGTCATCAAACTTAAATAATAATGTTCGAGGGAAAAAGACCTCTATGGCAT	530
Qy		GluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArgArgLysVal	140
Db		GAGTCTGATTTTGCTGTGGACAATGTTGGGAATAGAACTCTGCGCAGGAGGAGAAAGGTA	
Qy		LysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMetThrGlnPro	
Db		AAACGCATGGCAGTAGATCTCCCACAGGACATCTCTAACAAACGGACAATGACCCAGCCA	
Qy	0_	ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe	
Db		CCTGAGGGTTGTAGAGATCAGGACATGGACAGTGATAGAGCCTACCAGTATCAAGAATTT	
Qy Dh		ThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyProLysIleGln	
Db		ACCAAGAACAAAGTCAAAAAAAAGAAAGTTGAAAATCAGACAAGGACCAAAAATCCAA AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu	
Qy Db			
Qy		CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer	
Db			
Qy	241	SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer	260
Db			
Qy	261	AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp	280
Db	951	GACTGGTTCTACGAAAAGGAATCAGGTGGAGCATGTGGTATCACTGGAGTTGTGCCCTGG	1010
Qy	281	TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer	300
Db	1011	TGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTCTTTGAAAGT	1070
Qy	301	<pre>IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg</pre>	320
Db	1071	ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA	1130
Qy	321	LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr	340
Db	1131	CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAGGGACTCCAACT	1190
Qy	341	SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp	360
Db		TCÁÁTGGTÁCCCÁTTCCTGGCCCÁGTGGGTÁÁCÁÁGÁGÁÁTGGTTCÁTTTTTCCCCGGÁT	
Qy	361	SerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln	380

```
us-09-771-312-2.rng
Db
        1251 TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG 1310
         381 LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
Qy
        1311 CTTCTGAGAGATAATCGAGCTGAAAGAGGACACAAGAAAAATTGTTCTGTGAGAACAGCC 1370
Db
         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
             Db
        1371 AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA 1430
         421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
             Db
        1431 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1490
         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
             Db
        1491 ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA 1550
         461 GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys 480
Qy
Db
        1551 GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG 1610
         481 GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
             Db
        1611 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1670
         501 GlyLysserAla 504
Qy
             Db
        1671 GGAAAATCCGCC 1682
RESULT 3
ABN59701
    ABN59701 standard; cDNA; 2344 BP.
ΙD
XX
AC
    ABN59701;
XX
    28-JUN-2002 (first entry)
DT
XX
    Novel human coding sequence SEQ ID NO: 112.
DE
XX
    Human; antianaemic; vulnerary; antiinflammatory; immunomodulator; antiinfertility; cerebroprotective; cytostatic; rheumatic; gene therapy; neuroprotective; antiparkinsonian; protein therapy; EST;
KW
KW
KW
    expressed sequence tag; gene; ss.
KW
XX
os
    Homo sapiens.
XX
PΝ
    WO200222660-A2.
XX
PD
    21-MAR-2002.
XX
    10-SEP-2001; 2001WO-US026015.
PF
XX
PR
    11-SEP-2000; 2000US-00659671.
XX
    (HYSE-) HYSEQ INC.
PA
XX
ΡI
    Tang YT, Liu C, Zhou P, Asundi V, Zhang J, Zhao QA,
                                                            Ren F;
    Xue AJ, Yang Y, Wehrman T, Drmanac RT;
PΙ
XX
    WPI; 2002-292408/33.
DR
    P-PSDB; ABB97288.
DR
XX
```

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us-09-771-312-2.rng
PT
      An isolated polynucleotide for treating diseases associated with its
      encoded polypeptide such as cancer and multiple sclerosis.
PT
XX
PS
      Claim 1; SEQ ID NO 112; 509pp; English.
XX
      The present invention provides the protein and coding sequences of 444
CC
     The present invention provides the protein and coding sequences of 444 novel human proteins. These were isolated from expressed sequences tags (ESTs). They can be used to stimulate cell growth, to regulate haematopoiesis e.g. to treat aplastic anaemia, to help tissue regrowth e.g. in burn treatment, to regulate the immune system e.g. to treat multiple sclerosis, to regulate activin or inhibin e.g. to treat infertility, to regulate haemostasis or thrombolysis e.g. to treat stroke and cancer, to screen for drugs, to treat inflammatory conditions e.g. rheumatoid arthritis, and to treat nervous system disorders e.g. Parkinson's disease. The present sequence is a coding sequence of the invention
CC
      invention
XX
     Sequence 2344 BP; 747 A; 476 C; 541 G; 580 T; 0 U; 0 Other;
SQ
Alignment Scores:
                                                               2344
Pred. No.:
                            2.08e-199
                                              Length:
Score:
                            2694.00
                                                               504
                                              Matches:
Percent Similarity:
                            100.00%
                                              Conservative:
                                                               0
Best Local Similarity:
                            100.00%
                                              Mismatches:
                                                               0
Query Match:
                            100.00%
                                              Indels:
                                                               0
                                                               0
US-09-771-312-2 (1-504) x ABN59701 (1-2344)
Qy
              1 MetGluGluLeuValHisAspLeuValSerAlaLeuGluGluSerSerGluGlnAlaArg 20
                Db
           170 ATGGAGGAGCTGGTTCATGACCTTGTCTCAGCATTGGAAGAGCTCAGAGCAAGCTCGA 229
            21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40
Qy
Db
           230 GGTGGÁTTTGCTGÁÁÁCÁGGÁGÁCCÁTTCTCGÁÁGTÁTÁTCTTGCCCTCTGÁÁÁCGCCÁG 289
             41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60
Qy
           290 GCAAGGAAAAGGAGAGAGAAAACGGAGGTCGTATAATGTGCATCACCCGTGGGAGACT 349
nh
            61 GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg 80
Qу
Db
            350 GGTCACTGCTTAÁGTGAÁGGCTCTGÁTTCTÁGTTTAGAÁGAÁCCAÁGCAÁGGÁCTÁTÁGÁ 409
           Qy
Db
           101 LysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgProLeuTrpHis 120
Qy
                470 AAGCGCAGGCCGTCATCAAACTTAAATAATAATGTTCGAGGGAAAAGACCTCTATGGCAT 529
Db
           121 GluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArgArgLysVal 140
Qy
                530 GÁGTCTGÁTTTTGCTGTGGÁCÁATGTTGGGÁÁTÁGÁÁCTCTGCGCÁGGÁGGÁGÁÁÁGGTÁ 589
Db
Qy
           141 LysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMetThrGlnPro 160
                Db
           590 AAACGCATGGCAGTAGATCTCCCACAGGACATCTCTAACAAACGGACAATGACCCAGCCA 649
           161 ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe 180
Qy
```

		us-09-771-312-2.rng	
Db	650	CCTGAGGGTTGTAGAGATCAGGACATGGACAGTGATAGAGCCTACCAGTATCAAGAATTT	709
Qy		ThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyProLysIleGln	
Db		ACCAAGAACAAAGTCAAAAAAAAGAAAGTTGAAAATAATCAGACAAGGACCAAAAAATCCAA	
Qy		AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu	
Db		GATGAAGGAGTAGTTTTAGAAAGTGAGGAAACGAACCAGACCAATAAGGACAAAATGGAA	
Qy Db		CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer	
Qy		SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer	
Db			
Qy	261	AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp	280
Db	950		1009
Qy	281	TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer	300
Db	1010	TGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTCTTTGAAAGT	1069
Qy	301	<pre>IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg</pre>	320
Db		ATCTTAACTGGTTCTTTTCCCCTTATGTCÁCÁCCCÁÁGCÁGÁÁGÁGÁGÁTTTCCÁÁGCTÁGÁ	
Qy		LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr	
Db		CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAGGGACTCCAACT SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp	
Qy Db		TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTTCCCCGGAT	
Qy		SerHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln	
Db			
Qy	381	LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla	400
Db	1310		1369
Qy	401	SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg	420
Db	1370	AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA	1429
Qy	421	LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro	440
Db		AÁÁGCTGCÁCCTTTGCCTGGÁCCTÁCTÁCTGCÁGGÁTTTGTÁGGTGÁÁÁÁÁTGCCCÁGCCÁ	
Qy		<pre>IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer</pre>	
Db		ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA	
Qy Db		GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys	
Qy		GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla	

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us-09-771-312-2.rng
                        Db
               1610 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1669
                 501 GlyLysserAla 504
Qy
                         Db
               1670 GGAAAATCCGCC 1681
RESULT 4
AAS11663
        AAS11663 standard; cDNA; 2345 BP.
ID
XX
        AAS11663:
AC
XX
        24-OCT-2001 (first entry)
DT
XX
DE
        Prostate and testis-related gene 84P2A9 cDNA.
XX
        84P2A9; PCR primer; DNA adaptor; prostate; testis; tissue; cancer; ss; leukaemia; tumour; kidney; brain; bone; skin; ovary; breast; pancreas;
KW
KW
        colon; lung; cytostatic; gene therapy; antibody therapy; ribozyme; single chain monoclonal antibody; serum; blood; urine.
KW
KW
XX
        Homo sapiens.
05
XX
        WO200155391-A2.
PN
XX
PD
        02-AUG-2001.
XX
PF
        26-JAN-2001; 2001WO-US002651.
XX
PR
        26-JAN-2000; 2000US-0178560P.
XX
PA
         (UROG-) UROGENESYS INC.
XX
PΙ
         Jakobovits A, Afar DEH, Challita-Eid PM, Levin E, Mitchell SC;
PΙ
        Hubert RS;
XX
        WPI: 2001-502631/55.
DR
DR
        P-PSDB; AAU06524.
XX
PT
        New 84P2A9 gene and its encoded protein, useful for diagnosing and
        treating cancer, e.g. leukemia and cancer of the prostate, testis, kidney, brain or bone, or for eliciting an immune response.
PT
PT
XX
PS
        Claim 1; Fig 2; 149pp; English.
XX
        The nucleic acid sequences represent the 84P2A9 gene and the primers and adaptors used to amplify 84P2A9 DNA. 84P2A9 exhibits prostate and testis specific expression in normal adult tissue, but it is also aberrantly expressed in many cancers including leukaemia and tumours of the prostate, testis, kidney, brain, bone, skin, ovary, breast, pancreas, colon and lung. The 84P2A9 polynucleotide, its related protein and also pentide fragments of the protein are therefore useful for diagnosing and
CC
CC
CC
CC
CC
CC
         peptide fragments of the protein are therefore useful for diagnosing and
CC
        treating cancer. A vector comprising a polynucleotide which encodes a single chain monoclonal antibody, that immunospecifically binds to an 84P2A9-related protein, and a ribozyme capable of cleaving a polynucleotide having the 84P2A9 coding sequence, are both useful in the preparation of a composition for treating a patient with a cancer that expresses 84P2A9. The sequences can be used in diagnostic methods to monitor the level of 84P2A9 gene products in serum, blood, urine and
CC
CC
CC
CC
CC
CC
CC
CC
        tissue and to thereby detect the presence of cancerous cells
XX
SQ
         Sequence 2345 BP; 750 A; 476 C; 542 G; 577 T; 0 U; 0 Other;
```

Alignment Pred. No. Score: Percent S Best Loca Query Mat DB:	: imil l Si	arity:	2.08e-199 2694.00 100.00% 100.00% 100.00%	Length: Matches: Conservative: Mismatches: Indels: Gaps:	2345 504 0 0 0
us-09-771	-312	-2 (1-504)	x AAS11663 (1-	-2345)	
Qy	1	MetGluGlu	LeuValHisAspLeu	ValSerAlaLeuGlu	GluserserGluGlnAlaArg 20
Db	163	ATGGAGGAG	CTGGTTCATGACCTT	GTCTCAGCATTGGAA	
Qy	21	GlyGlyPhe	AlaGluThrGlyAsp	HisSerArgSerIle	SerCysProLeuLysArgGln 40
Db	223	GGTGGATTT	GCTGAAACAGGAGAC	CATTCTCGAAGTATA	
Qy	41	AlaArgLys	ArgArgGlyArgLys	ArgArgSerTyrAsn	ValHisHisProTrpGluThr 60
Db	283				GTGCATCACCCGTGGGAGACT 342
Qy	61	GlyHisCys	LeuSerGluGlySer	AspSerSerLeuGlu	GluProSerLysAspTyrArg 80
Db	343	GGTCACTGC	TTAAGTGAAGGCTCT	GATTCTAGTTTAGAA	GAACCAAGCAAGGACTATAGA 402
Qy	81				AspAspGlnMetLeuValAla 100
Db	403	GAGAATCAC	AATAATAATAAAAAA	AGATCACAGTGACTCT	GATGACCAAATGTTAGTAGCA 462
Qy	101	LysArgArg	ProSerSerAsnLeu	ıAsnAsnAsnValArg	GlyLysArgProLeuTrpHis 120
Db	463				GGGAAAAGACCTCTATGGCAT 522
Qy	121	GluSerAsp	PheAlaValAspAsr 	NalGlyAsnArgThr	LeuArgArgArgArgLysVal 140
Db	523	GAGTCTGAT	TTTGCTGTGGACAAT	GTTGGGAATAGAACT	CTGCGCAGGAGGAGAAAGGTA 582
Qy	141				LysArgThrMetThrGlnPro 160
Db	583	AAACGCATG	ĠĊĀĠŤĀĠĀŤĊŤĊĊĊĀ	CAGGACATCTCTAAC	AAACGGACAATGACCCAGCCA 642
Qy	161				AlaTyrGlnTyrGlnGluPhe 180
Db	643	ĊĊŢĠĂĠĠĠŢ	ŤĠŤĀĠĀĠĀŤĊĀĠĠĀĊ	CÁTGGÁCÁGTGÁTÁGÁ	GCCTACCAGTATCAAGAATTT 702
Qy	181	ThrLysAsn	LysValLysLysArg 	LysLeuLysIleIle	ArgGlnGlyProLysIleGln 200
Db	703	ACCAAGAAC	ÁÁÁĠŤĊÁÁÁÁÁÁÁÁÁ	AAGTTGAAAATAATC	AGACAAGGACCAAAAATCCAA 762
Qy	201				ThrAsnLysAspLysMetGlu 220
Db	763				ACCAATAAGGACAAAATGGAA 822
Qy	221	CysGluGlu	GlnLysValSerAsp 	oGluLeuMetSerGlu 	SerAspSerSerSerLeuSer 240
Db		TGTGAAGAG	CAAAAAGTCTCAGAT	GAGCTCATGAGTGAA	AGTGATTCCAGCAGTCTCAGC 882
Qy					GlnGlyAspAspGluGlnSer 260
Db	883	AGCACTGAT	GCTGGATTGTTTACC	:AATGATGAGGGAAGA	CAAGGTGATGATGAACAGAGT 942

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us-09-771-312-2.rng
        261 AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp 280
Qy
        943 GACTGGTTCTACGAAAAGGAATCAGGTGGAGCATGTGGTATCACTGGAGTTGTGCCCTGG 1002
Db
        281 TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer 300
Qy
Db
        1003 TGGGÁÁÁÁGGÁÁGÁTCCTÁCTGÁGCTÁGÁCÁÁÁÁÁTGTÁCCÁGÁTCCTGTCTTTGÁÁÁGT 1062
        301 IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg 320
Qy
            1063 ATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTCCAAGCTAGA 1122
Db
        321 LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr 340
Qy
Db
        1123 CTCAGTCGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAGGGACTCCAACT 1182
        341 SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp 360
Qy
            Db
        1183 TCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATTTTTCCCCGGAT 1242
        361 SerHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln 380
Qy
Db
        1243 TCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATGACCAGCATCAG 1302
        381 LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla 400
Qy
        1303 CTTCTGÁGÁGÁTAATCGAGCTGÁAÁGAGGÁCACAAGAÁAÁTTGTTCTGTGÁGAACAGCC 1362
Db
        401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
            1363 AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGAGATATCAAACGGAGAAGA 1422
Db
        421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
Db
        1423 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1482
        441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
            Db
        1483 ATCCTÁGÁÁÁÁTÁATATTGGÁÁÁÁCCGÁÁTGCTTCÁGÁÁTÁTGGGCTGGÁCGCCTGGGTCÁ 1542
       Qy
Db
        481 GlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
Db
        1603 GGÁTTÁGGÁCTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1662
        501 GlyLysSerAla 504
Qy
            1663 GGAAAATCCGCC 1674
Db
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ACN91982
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ID
XX
AC
    ACN91982;
XX
    02-DEC-2004 (first entry)
DT
XX
    Breast cancer related marker, seq id 13132.
DE
XX
    Cancer; breast; tumour; cytostatic; marker; detection; therapy; ds.
KW
```

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XX
      Homo sapiens.
os
XX
PΝ
      US2003099974-A1.
XX
PD
      29-MAY-2003.
XX
PF
      18-JUL-2002; 2002US-00198846.
XX
      18-JUL-2001; 2001US-0306220P.
PR
XX
      (MILL-) MILLENNIUM PHARM INC.
PA
XX
PΙ
      Lillie J, Xu Y, Wang Y, Steinmann K;
XX
      WPI; 2003-787014/74.
DR
XX
      Novel isolated polypeptide associated with breast cancer, useful for
PT
PT
      detecting presence of polypeptide in sample, as a marker for breast
PT
      cancer.
XX
      Disclosure; SEQ ID NO 13132; 36pp; English.
PS
XX
CC
      The invention relates to an isolated polypeptide (I) associated with
     breast cancer which is encoded by a nucleic acid molecule comprising a nucleotide sequence (S1). Further disclosed is an antibody that binds to the polypeptide of the invention. The activity of the polypeptide of the invention may be described as cytostatic. The antibody is useful for detecting the presence of (I) in a sample. Nucleic acid molecules of the
CC
CC
CC
CC
CC
      invention are useful in the detection of breast tumours. (I) is useful as
CC
      a marker for breast cancer and in breast cancer therapy. Sequences given
CC
      in records ACN78851-ACN92934 represent nucleic acid markers associated
CC
     with breast cancer. Note: The sequence listing does not form part of the specification but may be obtained in electronic format from the USPTO web site at seqdata.uspto.gov/sequence.html?DocID=20030099974
CC
CC
CC
XX
      Sequence 2583 BP; 813 A; 519 C; 575 G; 664 T; 0 U; 12 Other:
SQ
Alignment Scores:
                            2.35e-199
Pred. No.:
                                              Lenath:
                                                                2583
                            2694.00
Score:
                                              Matches:
                                                                504
Percent Similarity:
                            100.00%
                                              Conservative:
                                                                0
Best Local Similarity:
                            100.00%
                                              Mismatches:
                                                                0
Query Match:
                            100.00%
                                              Indels:
                                                                0
DB:
                                                                0
                            11
                                              Gaps:
US-09-771-312-2 (1-504) x ACN91982 (1-2583)
Qy
              1 MetGluGluLeuValHisAspLeuValSerAlaLeuGluGluSerSerGluGlnAlaArg 20
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            199 ÁTGGÁGGÁGCTGGTTCATGÁCCTTGTCTCÁGCÁTTGGÁÁGÁGÁGÁGCTCÁGÁGCAÁGCTCGÁ 258
             21 GlyGlyPheAlaGluThrGlyAspHisSerArgSerIleSerCysProLeuLysArgGln 40
Qy
Db
           259 GGTGGATTTGCTGAAACAGGAGACCATTCTCGAAGTATATCTTGCCCTCTGAAACGCCAG 318
             41 AlaArgLysArgArgGlyArgLysArgArgSerTyrAsnValHisHisProTrpGluThr 60
Qy
Db
            319 GCAAGGAAAAGGAGAGAGAAAACGGAGGTCGTATAATGTGCATCACCCGTGGGAGACT 378
             61 GlyHisCysLeuSerGluGlySerAspSerSerLeuGluGluProSerLysAspTyrArg 80
QУ
            Db
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Qy	81	GluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMetLeuValAla	100
Db	439		498
Qy	101	LysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgProLeuTrpHis	120
Db	499	AAGCGCAGGCCGTCATCAAACTTAAATAATAATGTTCGAGGGAAAAGACCTCTATGGCAT	558
Qy	121	GluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArgArgLysVal	140
Db	559	ĠĠĠŦĊŦĠĂŦŤŦŤĠĊŦĠŦĠĠÁĊĀĀŦĠŦŤĠĠĠĀĀŤĀĠĀĀĊŦĊŦĠĊĠĊĀĠĠĀĠĠĀĠĀĀĀĠĠŦĀ	618
Qy	141	LysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMetThrGlnPro	160
Db	619	ÀAACGCÀTGGCÀGTÀGÀTCTCCCÀCAGGÁCÁTCTCTÁACAAACGGÀCÁATGÀCCCÁGCCÁ	678
Qу		ProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyrGlnGluPhe	
Db		CCTGAGGGTTGTAGAGATCAGGACATGGACAGTGATAGAGCCTACCAGTATCAAGAATTT	
Qy		ThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyProLysIleGln	
Db		ACCAAGAACAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	
Qy		AspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAspLysMetGlu	
Db		GATGAAGGAGTAGTTTTAGAAAGTGAGGAAACGAACCAGACCAATAAGGACAAAATGGAA	
Qy		CysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSerSerLeuSer	
Db		TGTGAAGAGCAAAAAGTCTCAGATGAGCTCATGAGTGAAAGTGATTCCAGCAGTCTCAGC	
Qy		SerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAspGluGlnSer	
Db		AspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyValValProTrp	
Qy Db			
		TrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProValPheGluSer	
Qy Db			1098
Qy		IleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGlyPheGlnAlaArg	
Db			
Qy		LeuSerArgLeuHisGlyMetSerSerLysAsnIleLysLysSerGlyGlyThrProThr	
Db	1159		1218
Qy	341	SerMetValProIleProGlyProValGlyAsnLysArgMetValHisPheSerProAsp	360
Db	1219		1278
Qy	361	SerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisAspGlnHisGln	380
Db	1279		1338
Qy	381	LeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerValArgThrAla	400

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us-09-771-312-2.rng
        1339 CTTCTGAGAGATAATCGAGCTGAAAGAGGACACAAGAAAAATTGTTCTGTGAGAACAGCC 1398
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         401 SerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleLysArgArgArg 420
Qy
Db
        1399 AGCAGGCAAACAAGCATGCATTTAGGATCCTTATGCACGGGGGGATATCAAACGGAGAAGA 1458
          421 LysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluAsnAlaGlnPro 440
Qy
              1459 AAAGCTGCACCTTTGCCTGGACCTACTACTGCAGGATTTGTAGGTGAAAATGCCCAGCCA 1518
Db
         441 IleLeuGluAsnAsnIleGlyAsnArgMetLeuGlnAsnMetGlyTrpThrProGlySer 460
Qy
         1519 ATCCTAGAAAATAATATTGGAAACCGAATGCTTCAGAATATGGGCTGGACGCCTGGGTCA 1578
Db
          461 GlyLeuGlyArgAspGlyLysGlyIleSerGluProIleGlnAlaMetGlnArgProLys 480
Qy
              1579 GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG 1638
Db
         481 GlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla 500
Qy
Db
         1639 GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTACTACCCCCAATGCA 1698
          501 GlyLysSerAla 504
Qy
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Db
RESULT 6
AAS72189
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ID
XX
     AAS72189;
AC
XX
     13-FEB-2002 (first entry)
DT
XX
    DNA encoding novel human diagnostic protein #7993.
DE
XX
    Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW
     food supplement; medical imaging; diagnostic; genetic disorder; ss.
KW
XX
    Homo sapiens.
os
XX
    WO200175067-A2.
PΝ
XX
PD
     11-oct-2001.
XX
     30-MAR-2001; 2001WO-US008631.
PF
XX
     31-MAR-2000; 2000US-00540217.
23-AUG-2000; 2000US-00649167.
PR
PR
XX
PA
     (HYSE-) HYSEQ INC.
XX
PΙ
     Drmanac RT, Liu C, Tang YT;
XX
     WPI: 2001-639362/73.
DR
     P-PSDB; ABG08002.
DR
XX
     New isolated polynucleotide and encoded polypeptides, useful in
PT
     diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess
PT
PT
     biodiversity.
PT
XX
     Claim 1; SEQ ID NO 7993; 103pp; English.
PS
```

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XX
       The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal
CC
CC
CC
CC
CC
CC
        activity of (II) or to treat disease states involving (II). (II) is
CC
       useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and
CC
CC
CC
CC
CC
CC
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CC
       and to produce other types of data and products dependent on DNA and amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic coding sequences of the invention. Note: The sequence data for this patent did not appear in the printed specification, but was obtained in electronic format directly from WIPO at
CC
CC
cc
\mathsf{CC}
CC
        ftp.wipo.int/pub/published_pct_sequences
CC
XX
        Sequence 1563 BP; 486 A; 326 C; 395 G; 356 T; 0 U; 0 Other;
SQ
Alignment Scores:
Pred. No.:
                                     1.55e-106
1495.50
                                                             Length:
                                                                                    1563
Score:
                                                             Matches:
                                                                                    312
Percent Similarity:
                                     69.80%
                                                             Conservative:
                                     68.27%
Best Local Similarity:
                                                             Mismatches:
                                                                                    21
Query Match:
                                     55.51%
                                                             Indels:
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                                                             Gaps:
US-09-771-312-2 (1-504) x AAS72189 (1-1563)
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               155 ÁTGGÁGGÁGCTGGTTCÁTGÁCCTTGTCTCÁGCÁTTGGÁÁAGAGÁGCTCCAGAGCAAGCCT 214
Db
               Qy
Db
                 39 rgGlnAlaArgLysArgArgGlyArgLysArgArg-SerTyrAsnValHisHisProTrp 58
Qy
               275 GCCCAGCAAGGAAAAGGAGAGAGAGAAAACGGAGGTTCGTATAATGTGCATCACCCGTGG 334
Db
                 59 Glu-ThrGlyHisCysLeu--SerGluGlySerAspSerSerLeuGluGluProSerLys 77
Qy
               335 GÁGGÁCTGGTCÁCTGGCTTAAÁGTGÁÁGGCTCTGÁTTCTÁGT------ 376
Db
Qу
                 78 AspTyrArgGluAsnHisAsnAsnAsnLysLysAspHisSerAspSerAspAspGlnMet 97
               376 ----- 376
Db
                 98 LeuValAlaLysArgArgProSerSerAsnLeuAsnAsnAsnValArgGlyLysArgPro 117
Qy
               376 ----- 376
Db
               118 LeuTrpHisGluSerAspPheAlaValAspAsnValGlyAsnArgThrLeuArgArgArg 137
Qy
               376 ----- 376
Db
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		us-09-771-312-2.rng	
Qy		ArgLysValLysArgMetAlaValAspLeuProGlnAspIleSerAsnLysArgThrMet	
Db			
Qy		ThrGlnProProGluGlyCysArgAspGlnAspMetAspSerAspArgAlaTyrGlnTyr	
Db	376		376
Qy		GlnGluPheThrLysAsnLysValLysLysArgLysLeuLysIleIleArgGlnGlyPro	
Db	377	TTTAGAAGAACAÀAĠTCAÀAÀAÀAÀAĠAAĠTTĠAAAATAATĊAĠAĊAAĠĠAĊĊĀ	430
Qy	198	LysIleGlnAspGluGlyValValLeuGluSerGluGluThrAsnGlnThrAsnLysAsp	217
Db	431	AÁÁÁTCCÁÁGÁTGÁÁGGÁGTÁGTTTTÁGÁÁÁÁGTGÁGGÁÁACGÁÁCCÁGÁCCÁÁTÁÁGGÁC	490
Qy	218	LysMetGluCysGluGluGlnLysValSerAspGluLeuMetSerGluSerAspSerSer	237
Db	491	AÁÁÁTGGÁÁTGTGÁÁGÁGCÁÁÁÁÁÁGTCTCÁGÁTGÁGCTCÁTGÁGTGÁÁAGTGÁTTCCÁGC	550
Qy	238	SerLeuSerSerThrAspAlaGlyLeuPheThrAsnAspGluGlyArgGlnGlyAspAsp	257
Db		AGTCTCAGCAGCACTGATGCTGGATTGTTTACCAATGATGAGGGAAGACAAGGTGATGAT	
Qy	258	GluGlnSerAspTrpPheTyrGluLysGluSerGlyGlyAlaCysGlyIleThrGlyVal	277
Db	611	ĠĂĂĊĂĠĂĠŤĠĂĊŤĠĠŤŤĊŤĂĊĠĂĂĂĂĠĠĂĂŤĊĂĠĠŤĠĠĂĠĊĂŤĠŤĠĠŤĂŤĊĂĊŤĠĠĀĠŤŤ	670
Qy	278	ValProTrpTrpGluLysGluAspProThrGluLeuAspLysAsnValProAspProVal	297
Db		GTGCCTGGTGGGAAAAGGAAGATCCTACTGAGCTAGACAAAAATGTACCAGATCCTGTC	
Qy	298	PheGluSerIleLeuThrGlySerPheProLeuMetSerHisProSerArgArgGly-Ph	317
Db	731	TTTGAAAGTATCTTAACTGGTTCTTTTCCCCTTATGTCACACCCAAGCAGAAGAGGTTTT	790
Qy	317	eGlnAlaArgLeuSerArg-LeuHisGlyMetSerSerLysAsnIleLysLysSerGlyG	337
Db	791	CCAACTAAGACTCAGTCGGCCTTCATGGAATGTCTTCAAAGAATATTAAAAAATCTGGAG	850
Qy	337	<pre>lyThrProThrSerMetValProIleProGlyProValGlyAsnLysArgMetValHisP </pre>	357
Db		GGACTCCAACTTCAATGGTACCCATTCCTGGCCCAGTGGGTAACAAGAGAATGGTTCATT	
Qy	357	heSerProAspSerHisHisHisAspHisTrpPheSerProGlyAlaArgThrGluHisA	377
Db		TTTCCCCGGATTCTCATCACCATGACCATTGGTTTAGCCCTGGGGCTAGGACAGAGCATG	
Qy	377	<pre>spGlnHisGlnLeuLeuArgAspAsnArgAlaGluArgGlyHisLysLysAsnCysSerV</pre>	397
Db		ACCÁGCATCÁGCTTCTGÁGAGATÁATCGÁGCTGÁÁÁGÁGÁGÁCÁCÁÁGÁÁÁÁÁTTGTTCTG	
Qy	397	alArgThrAlaSerArgGlnThrSerMetHisLeuGlySerLeuCysThrGlyAspIleL	417
Db	1031	TGÁGÁÁCÁGCÁGCÁGGÁÁÁCÁÁGCÁTGCÁTTTÁGGÁTCCTTÁTGCÁCGGGÁGÁTÁTCÁ	1090
Qy	417	ysArgArgArgLysAlaAlaProLeuProGlyProThrThrAlaGlyPheValGlyGluA	437
Db	1091	AACGGAGAAGAAAGCTGCACCTTTGCCTGGACCTACTACTGCAGATTATTTCTCCCCCA	1150
Qy	437	snAlaGlnProIleLeuGluAsnAsnIleGlyAsn 448 ::: ::::: ::::	
Db	1151	TTCCCAAGCCAGTTATAGTAAAAGAATGTGGAAGT 1185	

GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: December 4, 2005, 10:07:40; Search time 41 Seconds

(without alignments)

1182.762 Million cell updates/sec

Title: US-09-771-312-2

Perfect score: 2694

Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5.

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

0.

Maximum Match 100%

Listing first 45 summaries

Database : PIR_80:*

1: pir1:*

2: pir2:*

3: pir3:*

4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

			₹				
Re	sult		Query				
_	No.	Score	Match	Length	DB	ID	Description
	1	177.5	6.6	1105	2	Т47582	hypothetical prote
	2	167.5	6.2	767	2	S63182	hypothetical prote
	3	156	5.8	542	2	T46464	hypothetical prote
	4	149	5.5	695	2	T40168	hypothetical prote
	5	148.5	5.5	1403	1	A47328	natural killer cel
	6	146.5	5.4	669	2	T28754	hypothetical prote
	7	143	5.3	1577	2	T19722	hypothetical prote
	8	143	5.3	3498	2	T22330	hypothetical prote
	9	138.5	5.1	368	2	G88636	protein W09G12.7 [
	10	135.5	5.0	643	2	A96636	unknown protein, 7
	11	134.5	5.0	699	2	I38073	nucleolar phosphop
	12	134.5	5.0	896	2	D96556	hypothetical prote
	13	133.5	5.0	1672	2	T46237	hypothetical prote

14	133	4.9	705	2	D88536	acidic protein - C
15	133	4.9	705	2	S27786	acidic protein - C
16	133	4.9	943	2	A42681	centromere protein
17	131.5	4.9	425	2	S55147	hypothetical prote
18	130	4.8	608	2	Т02299	hypothetical prote
19	130	4.8	679	2	S48437	hypothetical prote
20	129.5	4.8	2526	2	T20531	hypothetical prote
21	129.5	4.8	2722	2	T20532	hypothetical prote
22	129.5	4.8	2738	2	E88320	protein F07A11.6 [
23	128.5	4.8	543	2	T27190	hypothetical prote
24	128.5	4.8	552	2	Т27191	hypothetical prote
25	128.5	4.8	954	2	E86174	protein F19P19.26
26	127.5	4.7	493	2	т02376	hypothetical prote
27	127.5	4.7	539	2	T15256	hypothetical prote
28	127	4.7	763	2	Т08929	hypothetical prote
29	127	4.7	786	2	Т33856	hypothetical prote
30	127	4.7	845	2	A45669	neurofilament trip
31	127	4.7	963	2	T04002	hypothetical prote
32	126.5	4.7	390	2	Т34137	hypothetical prote
33	126	4.7	598	2	B40713	cylicin I - human
34	126	4.7	1032	2	A57514	RNA helicase HELll
35	125	4.6	1274	2	A89959	hypothetical prote
36	124.5	4.6	817	2	S53919	hypothetical prote
37	124	4.6	775	2	T21259	hypothetical prote
38	124	4.6	1166	2	Н86341	hypothetical prote
39	123.5	4.6	849	2	E86306	Similar to tufteli
40	123	4.6	529	2	Т50609	hypothetical prote
41	122	4.5	581	2	T22455	hypothetical prote
42	122	4.5	611	2	T22456	hypothetical prote
43	122	4.5	971	2	T24866	hypothetical prote
44	122	4.5	1230	2	T22458	hypothetical prote
45	121	4.5	4910	2	S64942	probable membrane

ALIGNMENTS

RESULT 1

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T47582
hypothetical protein F24B22.190 - Arabidopsis thaliana
C; Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 20-Apr-2000 #sequence revision 20-Apr-2000 #text change 09-Jul-2004
C; Accession: T47582
R; Bloecker, H.; Mewes, H.W.; Lemcke, K.; Mayer, K.F.X.; Quetier, F.; Salanoubat,
Μ.
submitted to the Protein Sequence Database, January 2000
A; Reference number: Z23016
A; Accession: T47582
A; Status: preliminary
A; Molecule type: DNA
A; Residues: 1-1105 <BLO>
A; Cross-references: UNIPROT: Q9M383; UNIPARC: UPI00000A410D; EMBL: AL132957
A; Experimental source: cultivar Columbia; BAC clone F24B22
C; Genetics:
A; Map position: 3
A; Introns: 35/3; 56/2; 294/3; 318/3; 349/3; 376/2; 426/3; 455/1; 485/3; 508/3;
568/3; 633/1; 662/3; 681/3; 710/2; 981/1; 1043/3
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A; Note: F24B22.190

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DŁ)	722	TQQYVPCPDQNNES-KVTENQPDSAKKEKSSQQKVIISAATTPNVEKVLSLPDAVQAAAA	780
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Qζ	7		NKRMVHFSPDSHHHDHWFSPGARTEHDQHQL-LRDNRAERGHKKNCSVRTASRQTSMH : : : : :	407
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Dk)	984	LMGLRKGSSDPTPFPPGVGGRGITTSTEVSSFDVITEERAIDESNVGNRMLRNM	103
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GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame plus p2n model Run on: December 11, 2005, 17:38:44; Search time 5807 Seconds (without alignments) 4060.735 Million cell updates/sec Title: US-09-771-312-2 Perfect score: 2694 Sequence: 1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504 Scoring table: BLOSUM62 Xgapop 10.0 , Xgapext 0.5 Ygapop 10.0 , Ygapext 0.5 Fgapop 6.0 , Fgapext 7.0 Delop 6.0 , Delext 7.0 41078325 segs, 23393541228 residues Searched: Total number of hits satisfying chosen parameters: 82156650 Minimum DB seq length: 0 Maximum DB seq length: 2000000000 Post-processing: Minimum Match 0% Maximum Match 100% Listing first 45 summaries Command line parameters: -MODEL=frame+ p2n.model -DEV=xlh Q=/cgn2 1/USPTO spool/US09771312/runat 01122005 145312 15071/app query.fasta 1.6 -DB=EST -QFMT=fastap -SUFFIX=rst -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pto -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000 -USER=US09771312 @CGN 1 1 5315 @runat 01122005 145312_15071 -NCPU=6 -ICPU=3 -NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG -DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7 Database : EST:* 1: gb est1:* 2: gb est2:* 3: gb est3:* 4: gb htc:* 5: gb est4:* 6: gb est5:* 7: qb est6:* 8: qb est7:*

> 9: gb_gss1:* 10: gb_gss2:*

11: gb_gss3:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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	4	2275	84.4	2746	4	AK029990	AK029990 Mus muscu
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ALIGNMENTS

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VERSION
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            Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
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            Hubisz, M.J., Fledel-Alon, A., Tanenbaum, D.M., Civello, D.,
            White, T.J., Sninsky, J.J., Adams, M.D. and Cargill, M.
            A Scan for Positively Selected Genes in the Genomes of Humans and
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            Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
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  JOURNAL
            Rockville, MD 20850, USA
            This sequence was made by sequencing genomic exons and ordering
COMMENT
            them based on alignment. Translation starts at the beginning of
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           Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
           Hubisz, M.J., Fledel-Alon, A., Tanenbaum, D.M., Civello, D.,
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           A Scan for Positively Selected Genes in the Genomes of Humans and
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           Nielsen, R., Bustamante, C., Clark, A.G., Glanowski, S., Sackton, T.B.,
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           White, T.J., Sninsky, J.J., Adams, M.D. and Cargill, M.
 TITLE
           Direct Submission
           Submitted (05-MAY-2005) Celera Genomics, 45 West Gude Drive,
 JOURNAL
           Rockville, MD 20850, USA
COMMENT
           This sequence was made by sequencing genomic exons and ordering
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Db	1453	GGCCTTGGACGAGATGGCAAGGGGATCTCTGAGCCAATTCAAGCCATGCAGAGGCCAAAG	1512
Qу	481	GlyLeuGlyLeuGlyPheProLeuProLysSerThrSerAlaThrThrThrProAsnAla	500
Db	1513	GGATTAGGACTTGGATTTCCTCTACCAAAAAGTACTTCCGCAACTGCTACCCCCAATGCA	1572
Qу	501	GlyLysSerAla 504	
Db	1573	GGAAAATCCGCC 1584	

us-09-771-312-2.rup

GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on:

December 4, 2005, 10:07:25; Search time 230 Seconds

(without alignments) 1546.027 Million cell updates/sec

Title:

US-09-771-312-2

Perfect score:

2694

Sequence:

1 MEELVHDLVSALEESSEQAR......GFPLPKSTSATTTPNAGKSA 504

Scoring table:

BLOSUM62

Gapop 10.0, Gapext 0.5

Searched:

2166443 seqs, 705528306 residues

Total number of hits satisfying chosen parameters:

2166443

Minimum DB seq length: 0 Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database:

UniProt_05.80:* 1: uniprot_sprot:*

2: uniprot_trembl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

			%				
Res	sult No.	Score	Query	Length	DB	ID	Description
	1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16	2694 2694 2310 2009.5 1813 1587.5 1538 1513.5 1283.5 1008 939.5 627.5 593.5 545.5 438 430 320.5	100.0 100.0 85.7 74.6 67.3 58.9 57.1 56.2 47.6 37.4 34.9 23.3 20.2 16.3 16.0	528 528 527 504 376 414 375 410 561 216 408 482 482 453 467 107 221	1 2 1 2 2 2 2 2 2 1 2 1 2 2 2 2 2 2 2 2	GPTC2_HUMAN Q5VYK7_HUMAN GPTC2_MOUSE Q5F3Y2_CHICK Q5VYK8_HUMAN Q4V7S5_XENLA Q9D3E7_MOUSE Q6AY15_RAT Q4RRB2_TETNG Q6PIX0_HUMAN Q5RJ37_BRARE CN118_MOUSE Q9H3M3_HUMAN CN118_HUMAN Q4RLV5_TETNG Q9CSX3_MOUSE Q9ULA8_HUMAN	Q9nw75 homo sapien Q5vyk7 homo sapien Q7tqc7 mus musculu Q5f3y2 gallus gall Q5vyk8 homo sapien Q4v7s5 xenopus lae Q9d3e7 mus musculu Q6ay15 rattus norv Q4rrb2 tetraodon n Q6pix0 homo sapien Q5rj37 brachydanio Q6pe65 mus musculu Q9h3m3 homo sapien Q9nwq4 homo sapien Q4rlv5 tetraodon n Q9csx3 mus musculu Q9ula8 homo sapien
	18 19	197 177.5	7.3 6.6	928 1007	2 2	Q6H4V9_ORYSA Q8VYR8_ARATH	Q6h4v9 oryza sativ Q8vyr8 arabidopsis
	20	177.5	6.6	1105	2	Q9M383_ARATH	Q9m383 arabidopsis
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us-09-771-312-2.rup
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732
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157.5
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                            853
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ALIGNMENTS

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05-JUL-2004 (Rel. 44, Last sequence update)
10-MAY-2005 (Rel. 47, Last annotation update)
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OX
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RC
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         Nakagawa S., Senoh A., Mizoguchi H., Goto Y., Shimizu F., Wakebe H.,
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Okumura K., Nagase T., Nomura N., Kikuchi H., Masuho Y., Yamashita R.,
Nakai K., Yada T., Nakamura Y., Ohara O., Isogai T., Sugano S.;
"Complete sequencing and characterization of 21,243 full-length human
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RT
         Nat. Genet. 36:40-45(2004).
RL
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Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
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         Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
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Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
"Generation and initial analysis of more than 15,000 full-length human
RA
RA
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         and mouse cDNA sequences."
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         Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
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CC
CC
                 Name=1;
                     IsoId=Q9NW75-1; Sequence=Displayed;
CC
\mathsf{CC}
                 Name=2;
CC
                     IsoId=Q9NW75-2; Sequence=VSP_010527, VSP_010528;
CC
                     Note=No experimental confirmation available:
CC
         -!- SIMILARITY: Contains 1 G-patch domain.
CC
         This Swiss-Prot entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC
CC
         the European Bioinformatics Institute. There are no restrictions on its use as long as its content is in no way modified and this statement is not
CC
CC
CC
         removed.
CC
         EMBL; AK001114; BAA91509.1; -; mRNA. EMBL; BC042193; AAH42193.1; -; mRNA. EMBL; BC063474; AAH63474.1; -; mRNA.
DR
DR
DR
         Ensembl; ENSG00000092978; Homo sapiens.
DR
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DR
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PROSITE; PS50174; G_PATCH; 1.
DR
DR
DR
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Alternative splicing.
467 513
                              us-09-771-312-2.rup
KW
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Missing (in isoform 2).
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               377
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D -> N (in Ref. 2; AAH42193).
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                        100.0%;
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                                                                Gaps
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Qy
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OC.
OC
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us-09-771-312-2.rup
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OX
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    Griffiths C.;
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EMBL; AC096641; CAH70664.1; JOINED; Genomic_DNA.
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    GO; GO:0005622; C:intracellular; IEA.
GO; GO:0003676; F:nucleic acid binding; IEA.
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05-JUL-2004 (Rel. 44, Created)
05-JUL-2005 (Rel. 44, Last sequence update)
13-SEP-2005 (Rel. 48, Last annotation update)
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GN
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us-09-771-312-2.rup
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OC.
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RC
                MEDLINE=22354683; PubMed=12466851; DOI=10.1038/nature01266;
RX
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Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
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RA
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                Birney E., Hayashizaki Y.;
                "Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs.";
RT
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RL
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RP
                NUCLEOTIDE SEQUENCE [LARGE SCALE MRNA] (ISOFORM 1).
               STRAIN=C57BL/6; TISSUE=Brain; MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
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               Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA
RA
RA
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              Hopkins R.F., Jordan H., Moore I., Max S.I., wang J., Hsien F., Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L., Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C., Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J., Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W., Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA
RA
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               Fahey J., Helton E., Ketteman M., Madan A., Rodrigues S., Sanchez A., Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA
RΑ
               Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
"Generation and initial analysis of more than 15,000 full-length human
RA
RA
RA
RA
RT
RT
                and mouse cDNA sequences.
                Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RL
                -!- ALTERNATIVE PRODUCTS:
\mathsf{CC}
                            Event=Alternative splicing; Named isoforms=3;
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           Note=No experimental confirmation available:
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CC
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           Note=No experimental confirmation available:
     -!- SIMILARITY: Contains 1 G-patch domain.
CC
CC
CC
     This Swiss-Prot entry is copyright. It is produced through a collaboration
     between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use as long as its content is in no way modified and this statement is not
CC
CC
CC
CC
     removed.
CC
     EMBL; AK030026; BAC26744.1; -; mRNA.
DR
     EMBL; AK053781; BAC35520.1; -; mRNA.

EMBL; AK083471; BAC38928.1; -; mRNA.

EMBL; BC054810; AAH54810.1; -; mRNA.

Ensembl; ENSMUSG00000039210; Mus musculus.
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     MGI; MGI:1915019; Gpatc2.
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SMART; SM00443; G_patch; 1.
PROSITE; PS50174; G_PATCH; 1.
Alternative splicing.
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FT
                                    E (in isoform 3).
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FT
                                   D -> Y (in Ref. 1; BAC26744).
S -> P (in Ref. 2).
                         251
FT
     CONFLICT
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                  367
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                          58218 MW; 4F4F29FA56BE06B7 CRC64;
     SEQUENCE
SQ
                         85.7%; Score 2310; DB 1; Length 527;
84.9%; Pred. No. 1.7e-130;
ative 35; Mismatches 41; Indels
  Query Match
  Best Local Similarity
 Matches 428; Conservative
                                                                    0;
                                                                        Gaps
                                                                                 0;
            1 MEELVHDLVSALEESSEQARGGFAETGDHSRSISCPLKRQARKRRGRKRRSYNVHHPWET 60
Qy
               24 MEELVHDLVSÁLÉESSEQÁRGGFÁÉTGEHSRNLSCPLKRQÁRKRRGRKRRSYNVHHPWÉT 83
Dh
           61 GHCLSEGSDSSLEEPSKDYRENHNNNKKDHSDSDDQMLVAKRRPSSNLNNNVRGKRPLWH 120
Qγ
               84 GHCLSEGSDSSLEEPSKDYREKHSNNKKDRSDSDDQMLVAKRRPSSNLSSSVRGKRLLWH 143
Db
          121 ESDFAVDNVGNRTLRRRRKVKRMAVDLPQDISNKRTMTQPPEGCRDQDMDSDRAYQYQEF 180
Qy
               144 ESDFAVDSLGNRTLRRRRKVKRMÁVDLPQDVSSKRTMTQLPEGCRÓQDMDNÓRÁSQYPEF 203
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          181 TKNKVKKRKLKIIRQGPKIQDEGVVLESEETNQTNKDKMECEEQKVSDELMSESDSSSLS 240
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               204 TRKKVKKKKLKGIRPGPKTQEEGGVLESEERSQPNKDRMEYEEQKASDELRSESDTSSLS 263
Db
Qy
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Qy	51 SHHHDHWFSPGARTEHDQHQLLRDNRAERGHKKNCSVRTASRQTSMHLGSLCTGDIKRRR 420
Db	:
Qy	21 KAAPLPGPTTAGFVGENAQPILENNIGNRMLQNMGWTPGSGLGRDGKGISEPIQAMQRPK 480
Db	
Qy	81 GLGLGFPLPKSTSATTTPNAGKSA 504 : : :
Db	04 GLGLGFPLPKSSPTSPAPTSGNPA 527